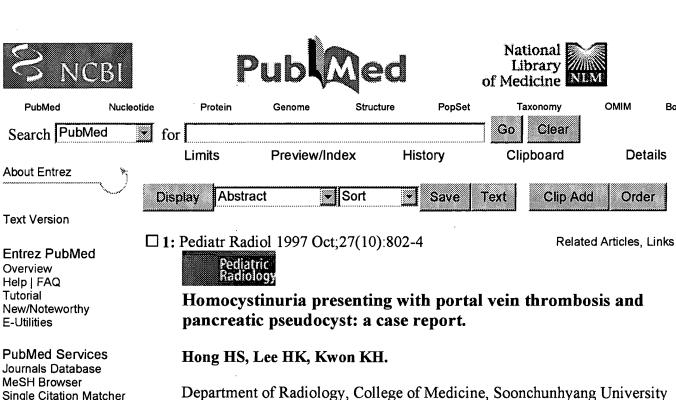
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Homocystinuria is a rare, inherited metabolic disease frequently associated with severe multisystemic involvement such as dislocated lenses, skeletal deformities, mental retardation, and premature vascular occlusion. Arterial and venous thromboembolic events present frequent and life-threatening complications in homocystinuric patients. It has been suggested that mild homocystinemia would be a risk factor for vascular disease.

PMID: 9323245 [PubMed - indexed for MEDLINE]

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